Under the Paperwork Reduction Act of 1995, no persons are required to

INFORMATION DISCLOSURE
STATEMENT BY APPLICANT
( Not for submission under 37 CFR 1.99)

Application Number		
Filing Date		
First Named Inventor	MCK	ENZIE, Duncan
Art Unit		
Examiner Name		
Attornov Docket Numb	or.	DDD 704440 IST

	U.S.PATENTS Remove										
Examiner Initial*	Cite No	Patent Number	Kind Code <sup>1</sup>	1 Issue Date		Name of Patentee or Applicant of cited Document		Pages,Columns,Lines when Relevant Passages or Relev Figures Appear			
	1										
If you wis	h to a	dd additional U.S. Pater	t citatio	n inform	ation pl	ease click the	Add button.		Add		
			U.S.P	ATENT	APPLI	CATION PUB	LICATIONS		Remove		
Examiner Initial*	Cite No	Publication Number	Kind Code <sup>1</sup>		Publication Date Name of Patentee or Applican of cited Document			Pages,Columns,Lines where Relevant Passages or Relevan Figures Appear			
	1										
If you wis	h to a	dd additional U.S. Publi	shed Ap	plication	citatio	n information p	lease click the Ad	d buttor	n. Add		
				FOREIG	SN PAT	TENT DOCUM	ENTS		Remove		
Examiner Initial*			Kind Code <sup>4</sup>	Publication Date	Applicant of cited		Pages,Columns,Lines where Relevant Passages or Relevant Figures Appear		74		
	1										
If you wish to add additional Foreign Patent Document citation information please click the Add button Add											
NON-PATENT LITERATURE DOCUMENTS Remove											
Examiner Initials*	Cite No	Include name of the au (book, magazine, journ	nal, seri	al, symp	osium,	catalog, etc), o					Тs

# Application Number | Filing Date | Find The Temperature | Find The T

	EYAMINED SIGNATURE								
If you wis	h to a	dd additional non-patent literature document citation information please click the Add button Add	_						
	4	M. GOMEZ-GALLEGO ET AL. "Synthesis of op-EDDHA and its detection as the main impurity in o.o-EDDHA commerical iron chelates", J. Agric. Food Chem., Vol. 50, 2002, Pages 6395-6399							
	3	L. HERNANDEZ-APAOLAZA ET AL: "Chromatographic determination of commercial Fe(III) chelates of ethylenediammetetraaceto acid, ethylenediammetel(o-hydroxyphenylacetic) acid and ethylenediamined(o-hydroxyphenylacetic) acid acid acid acid acid acid acid acid							
	2	M.A. CREMONINI ET AL: "NMR Analysis of the iron ligand ethylenediamined(io-hydroxyphenyljacetic acid (EDDHA) employed in fertilizers; J. Agric. Food Chern., Vol. 49, 2001, Pages 3527-3532							
	1	F. YUNTA ET AL: "Theoretical specification of ethylenediamine-N-(o-hydroxyphernylacetic)-N*-(p-hydroxyphernylacetic) acid (o.p-EDD+A) in agronomic conditions", J. Agric. Food Chem., Vol. 51, 2003, Pages 5391-5399							

Examiner Signature Date Considered

\*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 809. Draw line through a

citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

1 See Kind Code of USPTO Patent Documents at New USPTO\_GOU/ or MPEP 901.04. 2 Enter office that issued the document, by the how-left or potentially shaded \$7.5. 3 for the parents patent for comment, then chication of the parent of the paren

## Application Number Filing Date Filing Date STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99) Application Number MCXENZIE, Duncan Art Unit Examiner Name Attorney Cooket Number PDD 70444UST

### CERTIFICATION STATEMENT

Please see 37	CFR 1.97	and 1.9	38 to make the	appropriate	selection(s	s):
---------------	----------	---------	----------------	-------------	-------------	-----

That each item of information contained in the information disclosure statement was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of the information disclosure statement. See 37 CFR 197(eV1).

## OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquiry, no tem of information contained in the information disclosure statement was known to any individual designated in 37 CFR 1.58(c) more than three months prior to the filing of the information disclosure statement. Sex 7 CFR 1.97(c) in the filing of the information disclosure statement. Sex 7 CFR 1.97(c) in the filing of the information disclosure statement. Sex 7 CFR 1.97(c) in the filing of the information disclosure statement. Sex 7 CFR 1.97(c) in the filing of the information disclosure statement. Sex 7 CFR 1.97(c) in the filing of the information disclosure statement. Sex 9 CFR 1.97(c) in the filing of the information disclosure statement. Sex 9 CFR 1.97(c) in the filing of the information disclosure statement where the filing of the information disclosure statement.

- See attached certification statement.
- Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.
- .7 None

### SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

Signature	/William Teoli/	Date (YYYY-MM-DD)	2006-09-28
Name/Print	William A. Teoli, Jr.	Registration Number	33104

This collection of information is required by 37 CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is to file fand by the USPTO to process) an application. Confidentially is governed by \$5 U.S. C.12 and 37 CFR 1.14. This collection is estimated to take 1 hour to complete, including gathering, preparing and submitting the completed application from the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patient and Tradenant's Office, U.S. operationed for Commence, P. 0. Bot 1450, Alexandria, V.S. 2213.1-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, V.2.2313.1-1450.

## Privacy Act Statement

The Privacy Act of 1974 (P.L. 93-579) requires that you be given certain information in connection with your submission of the attached form related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, please be advised that. (1) the general authority for the collection of this information is 35 U.S.C. 2(b)(2); (2) famishing of the information solicided is voluntary, and (3) the principal purpose for which the information is used by the U.S. Patient and Trademan Kolfice is to process another examine your submission relation to a patient application or patient. If you do not furnish the requested process another examine your submission relation to the patient application or patient. If you do not furnish the requested the process another examines your submission, which may visually intermediate or for extension or about those when the basic high process another examines your submission, which may visually intermediate or for extension or a submission of the basic high process another examines your submission, which may

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
  - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement neodiations.
  - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record partains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
  - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974. as amended, pusuant to 5 U.S.C. 552a(m).
  - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
    may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
    to the Patent Cooperation Treaty.
  - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
- 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designe, cuting an inspection of records concluded by GSAs a part of that apency's responsibility to recommend improvements in records management practices and programs, under suthority of 4d U.S.C. 2004 and 2006. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make determinations about individuals.
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S. C. 122(b) or issuance of a patent pursuant to 35 U.S. C. 151. Further, a record may be disclosed, subject to the limitations of 37 CFR 1.14, as a routine use, to the public if the record via set float in an application which became abandomed or in which the proceedings were terminated and which application is referenced by either a published application, an application open to public inspections or an issuand patent.
  - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.